Progesterone for the Treatment of Traumatic Brain Injury (ProTECT III) Trial

Project Name: Phase III Clinical Trial to Determine if Progesterone Along With Standard Medical Care for Brain Injury is More Effective at Limiting the Amount of Damage Caused by a Traumatic Brain Injury Than Standard Medical Care Alone

Project Description: This study will determine the efficacy of administering intravenous (IV) progesterone (initiated within 4 hours of injury and administered for 72 hours, followed by an additional 24 hour taper) versus placebo for treating victims of moderate to severe acute TBI (Glasgow coma scale score 12-4).

Project Period Mar. 2010 – Jun. 2015

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Description of DCU Responsibilities: DCU is the Statistical and Data Management Center (SDMC), which provides statistical design and analysis, data management and IS infrastructure.